

**K010413 MINOLTA PULSOX-3SI, PULSOX-3IA, PULSOX-3LI**Aug 2, 2001  
171 days to decisionK010413 · Product code: **DQA** · AnesthesiologySource: <https://www.510kdatabase.net/k010413/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Oximeter (DQA)
Date received	Feb 12, 2001
Decision date	Aug 2, 2001
Days to decision	171 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

---

Company	<b>Konica Minolta Sensing, Inc.</b>
Location	Bailey, CO, US
Contact	NANCI DEXTER
510(k) history	3 submissions · 3 cleared · 2001-2006

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k010413/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated May 25, 2026